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M.L.
PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

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Group Art Unit: 1644

Examiner: Mary Tung

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For: POLYEPITOPE VACCINES

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PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Please amend the above-identified patent application as follows:

In the claims:

Please delete claims 1-¹³~~30~~ and add the following claims:

21.126 -- ¹⁴~~31~~. A polynucleotide comprising a nucleic acid sequence encoding at least two CTL epitopes,

wherein at least two of the epitopes are restricted by the same HLA gene.

AI 21.126
32. ¹⁵~~31~~. The polynucleotide of claim ¹⁴~~31~~, wherein the sequence encoding the CTL epitopes are contiguous.

R1.126 ¹⁶/₃₃ The polynucleotide of claim ¹⁴/₃₁, wherein said polynucleotide encodes two epitopes.

R1.126 ¹⁷/₃₄ The polynucleotide of claim ¹⁴/₃₁, wherein said polynucleotide encodes three epitopes.

R1.126 ¹⁸/₃₅ The polynucleotide of claim ¹⁴/₃₁, wherein said polynucleotide encodes nine epitopes.

R1.126 ¹⁹/₃₆ The polynucleotide of claim ¹⁴/₃₁, wherein said polynucleotide encodes ten epitopes.

R1.126 ²⁰/₃₇ The polynucleotide of claim ¹⁴/₃₁, wherein said polynucleotide encodes CTL epitopes from a plurality of pathogens.

R1.126 ²¹/₃₈ The polynucleotide of claim ¹⁴/₃₁, further defined as an expression vector.

R1.126 ²²/₃₉ The polynucleotide of claim ²¹/₃₈, wherein said vector is selected from the group consisting of a viral vector and a virus-like particle (VLP).

R1.126 ²³/₄₀ The polynucleotide of claim ²²/₃₉, wherein said viral vector is a vaccinia vector.

R1.126 ²⁴/₄₁ The polynucleotide of claim ²²/₃₉, wherein said viral vector is an avipox virus vector.

R1.126 ²⁵/₄₂ The polynucleotide of claim ²²/₃₉, wherein said vector is a VLP.

R1.126 ²⁶/₄₃ The polynucleotide of claim ¹⁴/₃₁, wherein at least one of said CTL epitopes is derived from a pathogen.

R1.126 ²⁷/₄₄ The polynucleotide of claim ¹⁴/₃₁, wherein said polynucleotide comprises a nucleic acid sequence encoding CTL epitopes derived from a plurality of pathogens.

R1.126 ²⁸/₄₅ The polynucleotide of claim ²⁶/₄₃, wherein said pathogen is selected from the group consisting of Epstein Barr Virus, Influenza Virus, Cytomegalovirus, Adenovirus and HIV.

R1.126 ²⁹/₄₀

The polynucleotide of claim ²⁷/~~44~~, wherein said pathogen is selected from the group consisting of Epstein Barr Virus, Influenza Virus, Cytomegalovirus, Adenovirus and HIV.

R1.126 ³⁴/₄₇

The polynucleotide of claim ¹⁴/~~31~~, wherein at least one of said epitopes is derived from a tumor protein.

R1.126 ³¹/₄₈

The polynucleotide of claim ¹⁴/~~31~~, further comprising a nucleic acid sequence encoding a T helper cell epitope, a B cell epitope, or a toxin.

R1.126 ³²/₄₉

The polynucleotide of claim ¹⁴/~~31~~, further comprising a nucleic acid sequence encoding a T helper cell epitope.

R1.126 ³³/₅₀

The polynucleotide of claim ¹⁴/~~31~~, further comprising a nucleic acid sequence encoding a B cell epitope.

R1.126 ³⁴/₅₁

The polynucleotide of claim ¹⁴/~~31~~, further comprising a nucleic acid sequence encoding a toxin.

R1.126 ³⁵/₅₂

A nucleic acid vaccine comprising a polynucleotide comprising a nucleic acid sequence encoding at least two CTL epitopes from one or more pathogens, wherein at least two of said epitopes are restricted by the same HLA gene, and an acceptable carrier.

R1.126 ³⁶/₅₃

A synthetic or recombinant protein comprising at least two CTL epitopes from one or more pathogens, wherein at least two of said epitopes are restricted by the same HLA gene.

R1.126 ³⁷/₅₄

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises two CTL epitopes.

R1.126 ³⁸/₅₅

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises three CTL epitopes.

R1.126 ³⁹/₅₆

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises nine CTL epitopes.

R1.126 ⁴⁰/₅₇

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises ten CTL epitopes.

R1.126 ⁴¹/₅₈

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises at least one CTL epitope derived from a pathogen.

A1.126 ⁴²/₅₉

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises CTL epitopes derived from a plurality of pathogens.

R1.126 ⁴³/₆₀

The synthetic or recombinant protein of claim ⁴¹/~~58~~, wherein said pathogen is selected from the group consisting of Epstein Barr Virus, Influenza Virus, Cytomegalovirus, Adenovirus and HIV.

R1.126 ⁴⁴/₆₁

The synthetic or recombinant protein of claim ⁴²/~~59~~, wherein said pathogen is selected from the group consisting of Epstein Barr Virus, Influenza Virus, Cytomegalovirus, Adenovirus and HIV.

R1.126 ⁴⁵/₆₂

The synthetic or recombinant protein of claim ³⁶/~~53~~, wherein said protein comprises at least one CTL epitopes from a tumor protein.

R1.126 ⁴⁶/₆₃

The synthetic or recombinant protein of claim ³⁶/~~53~~, further comprising a T helper cell epitope, a B cell epitope, or a toxin.

R1.126 ~~47.~~
64.

The synthetic or recombinant protein of claim ~~53~~³⁶, further comprising a T helper cell epitope.

R1.126 ~~48.~~
65.

The synthetic or recombinant protein of claim ~~53~~³⁶, further comprising a B cell epitope.

R1.126 ~~49.~~
66.

The synthetic or recombinant protein of claim ~~53~~³⁶, further comprising a toxin.

R1.126 ~~50.~~
67.

A polyepitope vaccine, the vaccine comprising a synthetic or recombinant protein comprising at least two CTL epitopes, wherein at least two of the epitopes are restricted by the same HLA gene.

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Conrad.
R1.126 ~~51.~~
68.

A method of vaccinating a subject against one or more pathogens which method comprises administering to the subject a polynucleotide comprising a nucleic acid sequence encoding at least two CTL epitopes, wherein at least two of the epitopes are restricted by the same HLA gene.

R1.126 ~~52.~~
69.

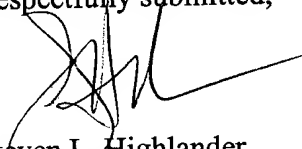
A method of vaccinating a subject against one or more pathogens which method comprises administering to the subject a synthetic or recombinant protein comprising at least two CTL epitopes, wherein at least two of the epitopes are restricted by the same HLA gene. --

A fee as set forth in 37 C.F.R. §§ 1.16-1.21 in the amount of \$1266.00 is enclosed herewith.

If an appropriate check has not been enclosed, or if it is insufficient under 37 C.F.R §§ 1.16 to 1.21, the Commissioner is hereby authorized to deduct any necessary fees from Fulbright & Jaworski Deposit Account No. 50-1212/10011879/01973.

Should Examiner Tung have any questions regarding this communication, she is invited to contact the undersigned at the telephone number listed below.

Respectfully submitted,



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